## AMENDMENTS TO THE CLAIMS

## 1. - 3. (Canceled)

- 4. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a percent time in mode switch source.
- 5. (Withdrawn) The method of claim 1, wherein the at least one additional source includes an R-wave and P-wave amplitude source.
- 6. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a reversion pace count source.
- 7. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a refractory sense count source.
- 8. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a high rate episode count source.
- 9. (Withdrawn) The method of claim 1, wherein the at least one additional source includes a time from implant source.
- 10. (Canceled)
- 11. (Withdrawn) The method of claim 2, wherein the message indicates a lead conductor or connector issue.
- 12. (Withdrawn) The method of claim 2, wherein the message indicates a lead insulation issue.

## 13. - 14. (Canceled)

15. (Withdrawn) The method of claim 13, wherein the biological interface issue includes lead dislodgement.

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- 16. (Withdrawn) The method of claim 13, wherein the biological interface issue includes exit block.
- 17. (Currently Amended) A method of lead status monitoring in an implantable medical device (IMD) comprising the steps of:

collecting lead impedance data,

collecting stimulation threshold data,

sensing signals along two distinct sensing pathways;

collecting data relating to one of a percent of time in mode switch, R-wave amplitude, P-wave amplitude, reversion pace count, refractory sense count, high rate episode count, and time from implant; and

processing the collected data in accordance with an algorithm having an integrated set of rules to determine if a lead status event has occurred, wherein each rule of the set applies a specific determination criterion to a particular aspect of the collected data, and wherein a first determination criterion is applied for signals sensed along a first sensing pathway of the two distinct sensing pathways, and a second determination criterion is applied for signals sensed along a second sensing pathway of the two distinct sensing pathways.

- 18. (Previously Presented) The method of claim 17, further comprising providing a message indicating a lead-related condition to a user based on the lead status event.
- 19. (Previously Presented) The method of claim 18, wherein the message indicates one of a lead conductor or connector issue, a lead insulation issue, and a biological interface issue.
- 20. (Previously Presented) The method of claim 19, wherein the biological interface issue includes one of myocardial perforation, lead dislodgement, and exit block.

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- 21. (Previously Presented) The method of claim 17, wherein the processing comprises: assigning weighted values to the collected data sets; and summing the assigned weighted values to determine if one of a plurality of lead status events has occurred.
- 22. (New) A method of distinguishing lead-related conditions in a medical device, comprising:

sensing signals received by a plurality of electrodes positioned along one or more leads;

determining whether a number of sensed events occurring along a first sensing pathway formed by one or more of the plurality of electrodes is greater than a first threshold associated with the first sensing pathway to generate a first event count;

determining whether a number of sensed events occurring along a second sensing pathway formed by one or more of the plurality of electrodes, different from the first sensing pathway, is greater than a second threshold associated with the second sensing pathway to generate a second event count; and

identifying the presence of a lead-related condition in response to the first event count and the second event count.

- 23. (New) The method of claim 22, wherein the first sensing pathway corresponds to a unipolar sensing pathway and the second sensing pathway corresponds to a bipolar sensing pathway.
- 24. (New) The method of claim 22, further comprising:

determining a number of counter windows; and

determining whether a sum of the first event count and the second event count is greater than a count threshold, wherein the count threshold varies depending on the determined number of counter windows.